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NEWS  3   FEB 28  PATDPAFULL - New display fields provide for legal status
                  data from INPADOC
NEWS  4   FEB 28  BABS - Current-awareness alerts (SDIs) available
NEWS  5   MAR 02  GBFULL: New full-text patent database on STN
NEWS  6   MAR 03  REGISTRY/ZREGISTRY - Sequence annotations enhanced
NEWS  7   MAR 03  MEDLINE file segment of TOXCENTER reloaded
NEWS  8   MAR 22  KOREAPAT now updated monthly; patent information enhanced
NEWS  9   MAR 22  Original IDE display format returns to REGISTRY/ZREGISTRY
NEWS  10  MAR 22  PATDPASPC - New patent database available
NEWS  11  MAR 22  REGISTRY/ZREGISTRY enhanced with experimental property tags
NEWS  12  APR 04  EPFULL enhanced with additional patent information and new
                  fields
NEWS  13  APR 04  EMBASE - Database reloaded and enhanced
NEWS  14  APR 18  New CAS Information Use Policies available online
NEWS  15  APR 25  Patent searching, including current-awareness alerts (SDIs),
                  based on application date in CA/CAplus and USPATFULL/USPAT2
                  may be affected by a change in filing date for U.S.
                  applications.
NEWS  16  APR 28  Improved searching of U.S. Patent Classifications for
                  U.S. patent records in CA/CAplus
NEWS  17  MAY 23  GBFULL enhanced with patent drawing images
NEWS  18  MAY 23  REGISTRY has been enhanced with source information from
                  CHEMCATS
NEWS  19  JUN 06  The Analysis Edition of STN Express with Discover!
                  (Version 8.0 for Windows) now available
NEWS  20  JUN 13  RUSSIAPAT: New full-text patent database on STN
NEWS  21  JUN 13  FRFULL enhanced with patent drawing images
NEWS  22  JUN 27  MARPAT displays enhanced with expanded G-group definitions
                  and text labels
NEWS  23  JUL 01  MEDICONF removed from STN
NEWS  24  JUL 07  STN Patent Forums to be held in July 2005
NEWS  25  JUL 13  SCISEARCH reloaded
NEWS  26  JUL 20  Powerful new interactive analysis and visualization software,
                  STN AnaVist, now available

NEWS EXPRESS  JUNE 13 CURRENT WINDOWS VERSION IS V8.0, CURRENT
              MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005

NEWS HOURS    STN Operating Hours Plus Help Desk Availability
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NEWS WWW      CAS World Wide Web Site (general information)

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FILE 'HOME' ENTERED AT 17:59:30 ON 23 JUL 2005

=> file reg
COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
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FULL ESTIMATED COST 0.21 0.21

FILE 'REGISTRY' ENTERED AT 17:59:42 ON 23 JUL 2005

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 22 JUL 2005 HIGHEST RN 856698-04-9
DICTIONARY FILE UPDATES: 22 JUL 2005 HIGHEST RN 856698-04-9

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TSCA INFORMATION NOW CURRENT THROUGH JANUARY 18, 2005

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*                                                                    *
* The CA roles and document type information have been removed from  *
* the IDE default display format and the ED field has been added,    *
* effective March 20, 2005.  A new display format, IDERL, is now     *
* available and contains the CA role and document type information.  *
*                                                                    *
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Structure search iteration limits have been increased. See HELP SLIMITS for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at:
http://www.cas.org/ONLINE/DBSS/registryss.html

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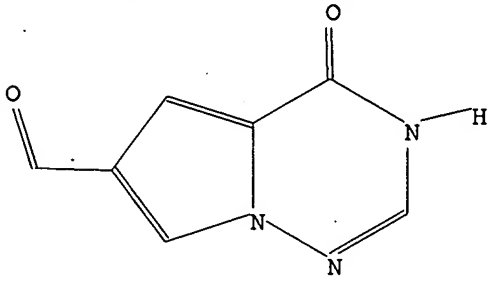

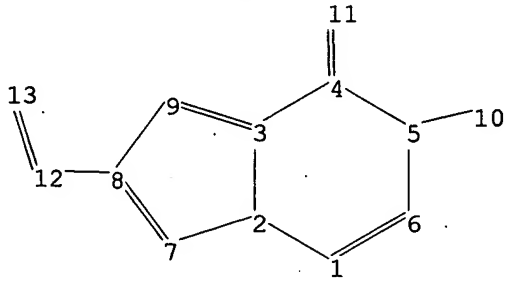

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10 11 12 13
ring nodes :
1 2 3 4 5 6 7 8 9
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ring bonds :
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exact/norm bonds :
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exact bonds :

10/773,002

5-10 8-12

Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS
11:CLASS 12:CLASS 13:CLASS

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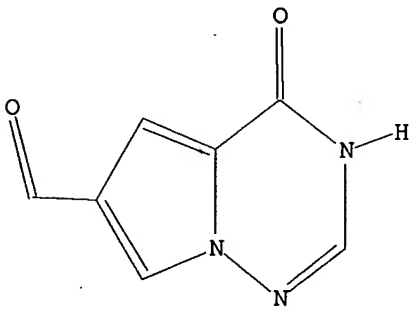

Structure attributes must be viewed using STN Express query preparation.

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SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 56 TO 504
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 sss ful
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FULL SCREEN SEARCH COMPLETED - 487 TO ITERATE

100.0% PROCESSED 487 ITERATIONS 23 ANSWERS
SEARCH TIME: 00.00.01

L3 23 SEA SSS FUL L1

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COST IN U.S. DOLLARS SINCE FILE ENTRY TOTAL SESSION
FULL ESTIMATED COST 161.33 161.54

FILE 'CAPLUS' ENTERED AT 18:00:20 ON 23 JUL 2005

FILE COVERS 1907 - 23 Jul 2005 VOL 143 ISS 5
FILE LAST UPDATED: 22 Jul 2005 (20050722/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l3
L4 15 L3

=> d l4 1-15 bib hitstr

L4 ANSWER 1 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2005:567104 CAPLUS
TI A preparation of (indazolylamino)pyrrolo[2,1-f][1,2,4]triazine derivatives, useful as antiproliferative agents
IN Swaminathan, Shankar; Gavai, Ashvinikumar V.; Fan, Junying; Patel, Bharat P.; Norris, Derek J.; Corbett, Richard Michael; Zheng, Bin
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 51 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005058245	A2	20050630	WO 2004-US41920	20041210
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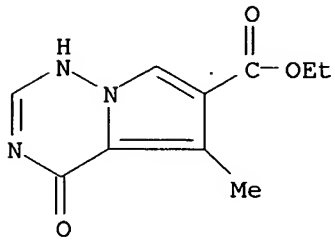
	RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRAI	US 2003-529347P	P	20031212		
	US 2004-8719	A	20041209		

IT **427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of (indazolylamino)pyrrolo[2,1-f][1,2,4]triazine derivs. useful as antiproliferative agents)
RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 2 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2005:160831 CAPLUS
DN 142:261564
TI Heteroaryl-substituted pyrrolo-triazine compounds useful as kinase inhibitors, particularly p38 kinases, and their preparation, pharmaceutical compositions, and use
IN Leftheris, Katerina; Wrobleski, Stephen T.; Dyckman, Alaric J.
PA USA
SO U.S. Pat. Appl. Publ., 29 pp., Cont.-in-part of U.S. Ser. No. 420,399.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2005043306	A1	20050224	US 2003-678388	20031003
	US 2004082582	A1	20040429	US 2003-420399	20030422
	WO 2005037838	A1	20050428	WO 2004-US30829	20040921

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRAI	US 2002-374938P	P	20020423
	US 2003-420399	A2	20030422
	US 2003-678388	A	20031003

OS MARPAT 142:261564
IT **621685-55-0P 621685-56-1P 621685-58-3P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of heteroaryl pyrrolotriazine compds. as p38 kinase inhibitors)
RN 621685-55-0 CAPLUS
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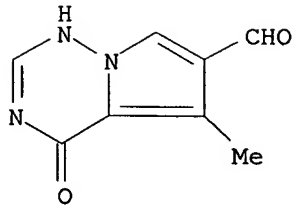

RN 621685-56-1 CAPLUS
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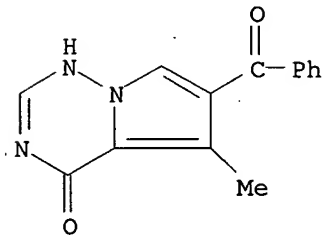

RN 621685-58-3 CAPLUS
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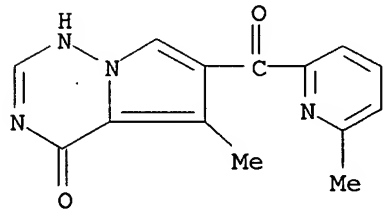
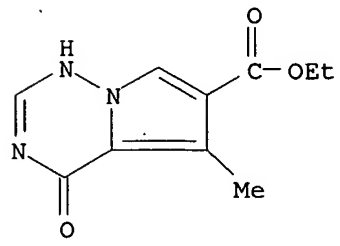
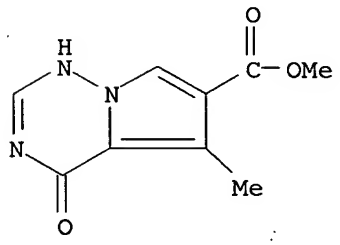


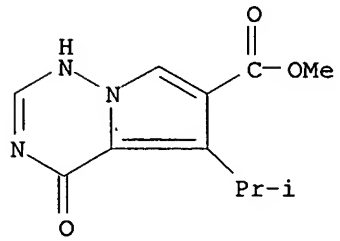
IT **427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of heteroaryl pyrrolotriazine compds. as p38 kinase inhibitors)
RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



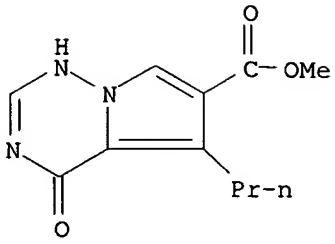
L4 ANSWER 3 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2005:130304 CAPLUS
DN 142:392380
TI Synthesis and SAR of 4-(3-hydroxyphenylamino)pyrrolo[2,1-f][1,2,4]triazine based VEGFR-2 kinase inhibitors
AU Borzilleri, Robert M.; Cai, Zhen-Wei; Ellis, Christopher; Fargnoli, Joseph; Fura, Aberra; Gerhardt, Tracy; Goyal, Bindu; Hunt, John T.; Mortillo, Steven; Qian, Ligang; Tokarski, John; Vyas, Viral; Wautlet, Barri; Zheng, Xioping; Bhide, Rajeev S.
CS Departments of Oncology Chemistry, Bristol-Myers Squibb Pharmaceutical Research Institute, Princeton, NJ, 08543-4000, USA
SO Bioorganic & Medicinal Chemistry Letters (2005), 15(5), 1429-1433
CODEN: BMCLE8; ISSN: 0960-894X
PB Elsevier B.V.
DT Journal
LA English
IT **310431-29-9P 651744-51-3P 850085-64-2P**
850085-66-4P 850085-97-1P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(synthesis and SAR of 4-(3-hydroxyphenylamino)pyrrolo[2,1-f][1,2,4]triazine based VEGFR-2 kinase inhibitors)
RN 310431-29-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



RN 651744-51-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



RN 850085-64-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-4-oxo-5-propyl-, methyl ester (9CI) (CA INDEX NAME)



RN 850085-66-4 CAPLUS
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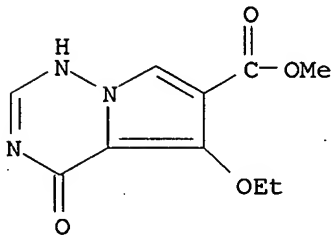
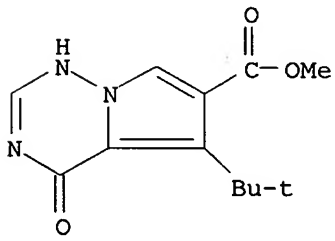


RN 850085-97-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-(1,1-dimethylethyl)-1,4-dihydro-4-oxo-, methyl ester (9CI) (CA INDEX NAME)

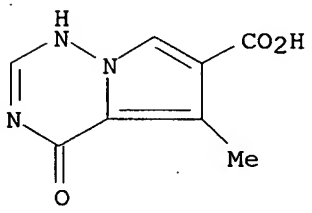


RE.CNT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

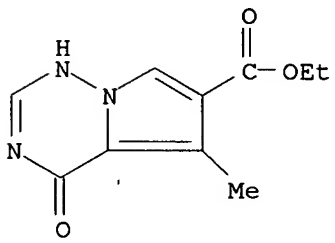
L4 ANSWER 4 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:654775 CAPLUS
DN 141:190807
TI Process for preparing pyrrolotriazine kinase inhibitors
IN Chen, Bang-Chi; Zhao, Rulin; Sundeen, Joseph Edward; Leftheris, Katerina; Hynes, John; Wrobleski, Stephen T.
PA USA
SO U.S. Pat. Appl. Publ., 20 pp.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

PI US 2004157846 A1 20040812 US 2004-773002 20040205
WO 2004072030 A2 20040826 WO 2004-US3223 20040205
WO 2004072030 A3 20041028
W: AE, AE, AG, AL, AL, AM, AM, AM, AT, AT, AU, AZ, AZ, BA, BB, BG, BG, BR, BR, BW, BY, BY, BZ, BZ, CA, CH, CN, CN, CO, CO, CR, CR, CU, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EC, EC, EE, EE, EG, ES, ES, FI, FI, GB, GD, GE, GE, GH, GM, HR, HR, HU, HU, ID, IL, IN, IS, JP, JP, KE, KE, KG, KG, KP, KP, KP, KR, KR, KZ, KZ, KZ, LC, LK, LR, LS, LS, LT, LU, LV, MA, MD, MD, MG, MK, MN, MW, MX, MX, MZ, MZ, NA, NI
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2003-445224P P 20030205
OS CASREACT 141:190807; MARPAT 141:190807
IT **310435-15-5P 427878-70-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(process for preparing pyrrolotriazine p38 kinase inhibitors)
RN 310435-15-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 5 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:531311 CAPLUS
DN 141:89122
TI Preparation of C-6 modified indazolyl pyrrolotriazines as antiproliferative agents
IN Vite, Gregory D.; Gavai, Ashvinikumar V.; Fink, Brian E.; Mastalerz, Harold; Kadow, John F.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 81 pp.

CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004054514	A2	20040701	WO 2003-US39542	20031212
WO 2004054514	A3	20041007		

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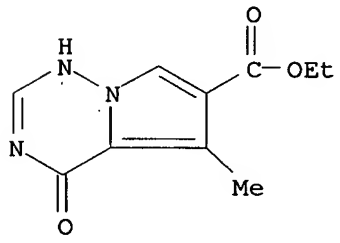
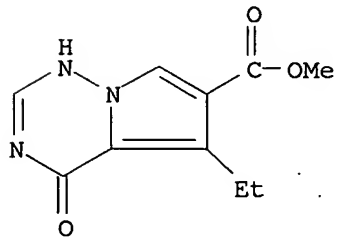
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2004142931	A1	20040722	US 2003-736476	20031215
US 6916815	B2	20050712		
PRAI US 2002-433190P	P	20021213		

OS MARPAT 141:89122
IT **427878-70-4 714971-30-9**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of C-6 modified indazolyl pyrrolotriazines as antiproliferative agents)
RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RN 714971-30-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-ethyl-1,4-dihydro-4-oxo-, methyl ester (9CI) (CA INDEX NAME)

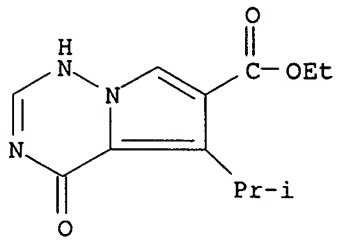


L4 ANSWER 6 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:120859 CAPLUS

DN 140:181471
TI Preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors for the treatment of cancer
IN Bhide, Rajeev S.; Borzilleri, Robert M.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

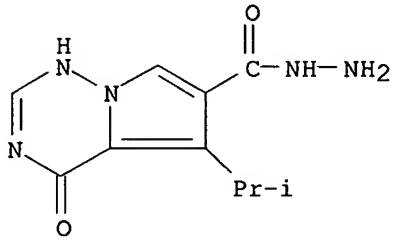
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2004013145	A1	20040212	WO 2003-US24273	20030804
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
US 2004063708	A1	20040401	US 2003-633997	20030804
EP 1543009	A1	20050622	EP 2003-767116	20030804
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
PRAI US 2002-400572P	P	20020802		
WO 2003-US24273	W	20030804		

OS MARPAT 140:181471
IT **651744-40-0P**
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors)
RN 651744-40-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)

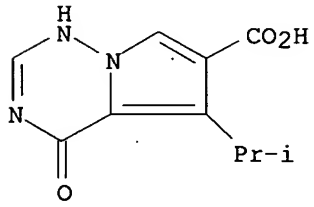


IT **658085-53-1P 658085-61-1P 658085-62-2P**
658085-63-3P 658085-64-4P 658085-69-9P
658085-71-3P, 4-Hydroxy-5-(1-methylethyl)pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid(2-oxopropyl)amide
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of pyrrolotriazines as tyrosine kinase activity inhibitors of growth factor receptors)
RN 658085-53-1 CAPLUS

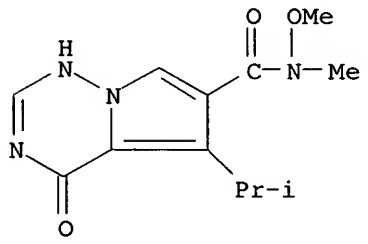
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, hydrazide (9CI) (CA INDEX NAME)



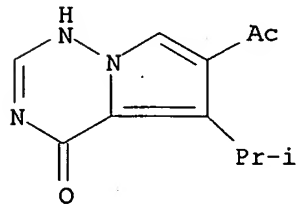
RN 658085-61-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo- (9CI) (CA INDEX NAME)



RN 658085-62-2 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-N-methoxy-N-methyl-5-(1-methylethyl)-4-oxo- (9CI) (CA INDEX NAME)

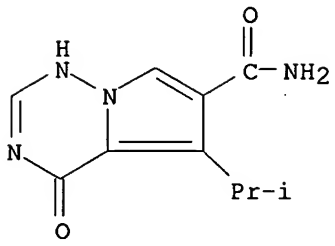


RN 658085-63-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-acetyl-5-(1-methylethyl)- (9CI) (CA INDEX NAME)

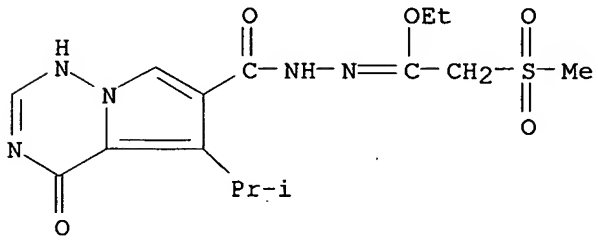


RN 658085-64-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-5-(1-methylethyl)-

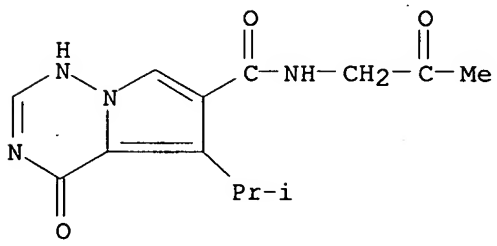
4-oxo- (9CI) (CA INDEX NAME)



RN 658085-69-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, [1-ethoxy-2-(methylsulfonyl)ethylidene]hydrazide (9CI) (CA INDEX NAME)



RN 658085-71-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxamide, 1,4-dihydro-5-(1-methylethyl)-4-oxo-N-(2-oxopropyl)- (9CI) (CA INDEX NAME)



L4 ANSWER 7 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:80847 CAPLUS
DN 140:124558
TI Pyrrolotriazine inhibitors of kinases for use in treatment of diseases associated with growth factor receptor signal transduction
IN Bhide, Rajeev; Cai, Zhen-wei; Qian, Ligang; Barbosa, Stephanie
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 84 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 3

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

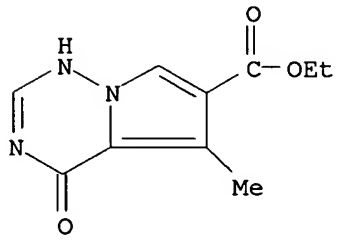
PI	WO 2004009784	A2	20040129	WO 2003-US22826	20030718
	WO 2004009784	A3	20040422		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2492804	AA	20040129	CA 2003-2492804	20030718
	US 2004063707	A1	20040401	US 2003-622593	20030718
	US 2004072832	A1	20040415	US 2003-623171	20030718
	US 6869952	B2	20050322		
	EP 1534290	A2	20050601	EP 2003-765881	20030718
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	US 2005124621	A1	20050609	US 2005-35248	20050113
PRAI	US 2002-397256P	P	20020719		
	US 2003-447213P	P	20030213		
	US 2003-623171	A1	20030718		
	WO 2003-US22826	W	20030718		

OS MARPAT 140:124558

IT **427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(pyrrolotriazine inhibitors of kinases for use in treatment of diseases associated with growth factor receptor signal transduction)

RN 427878-70-4 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 8 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:80698 CAPLUS

DN 140:146173

TI Preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases

IN Bhide, Rajeev; Ruel, Rejean; Thibeault, Carl; L'heureux, Alexandre

PA Bristol-Myers Squibb Company, USA

SO PCT Int. Appl., 66 pp.
CODEN: PIXXD2

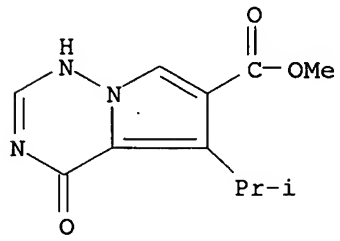
DT Patent

LA English

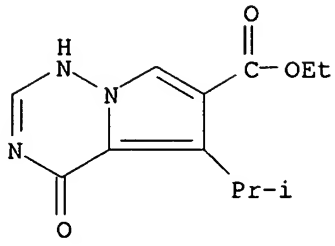
FAN.CNT 3

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

PI WO 2004009601 A1 20040129 WO 2003-US22554 20030718
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
CA 2492665 AA 20040129 CA 2003-2492665 20030718
US 2004063707 A1 20040401 US 2003-622593 20030718
US 2004072832 A1 20040415 US 2003-623171 20030718
US 6869952 B2 20050322
EP 1539763 A1 20050615 EP 2003-765754 20030718
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
US 2005124621 A1 20050609 US 2005-35248 20050113
PRAI US 2002-397256P P 20020719
US 2003-447213P P 20030213
US 2003-623171 A1 20030718
WO 2003-US22554 W 20030718
OS MARPAT 140:146173
IT **651744-51-3**
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases)
RN 651744-51-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



IT **651744-40-0P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as selective VEGFR-2 and FGFR-1 kinase inhibitors for treatment of proliferative diseases)
RN 651744-40-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-(1-methylethyl)-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)

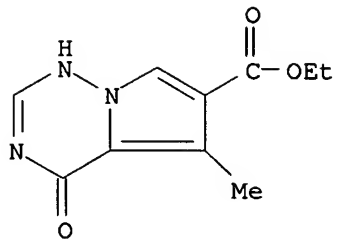


RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 9 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2004:80644 CAPLUS
DN 140:146018
TI Process for preparation of indolyloxypyrrolotriazines and their use as drugs.
IN Bhide, Rajeev; Fan, Junying; Parlanti, Luca; Barbosa, Stephanie; Qian, Ligang; Cai, Zhen-wei; Gibson, Francis S.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 48 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 3

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004009542	A2	20040129	WO 2003-US22755	20030721
	WO 2004009542	A3	20040513		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 2004077858	A1	20040422	US 2003-622280	20030718
	CA 2492861	AA	20040129	CA 2003-2492861	20030721
	EP 1554281	A2	20050720	EP 2003-765846	20030721
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	US 2005124621	A1	20050609	US 2005-35248	20050113
PRAI	US 2002-397256P	P	20020719		
	US 2003-447213P	P	20030213		
	US 2003-622280	A	20030718		
	US 2003-623171	A1	20030718		
	WO 2003-US22755	W	20030721		

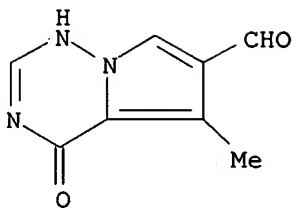
OS MARPAT 140:146018
IT **427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(process for preparation of indolyloxypyrrolotriazines and their use as drugs)
RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



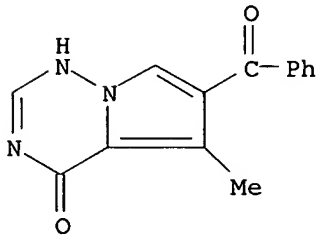
L4 ANSWER 10 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:875265 CAPLUS
DN 139:364963
TI Aryl ketone pyrrolo-triazine compounds useful as kinase inhibitors, particularly p38 kinases, and their preparation, pharmaceutical compositions, and use
IN Dyckman, Alaric; Leftheris, Katerina; Hynes, John
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 45 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003091229	A1	20031106	WO 2003-US12420	20030418
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	EP 1503996	A1	20050209	EP 2003-718493	20030418
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	US 2003232831	A1	20031218	US 2003-420445	20030422
PRAI	US 2002-374907P	P	20020423		
	WO 2003-US12420	W	20030418		

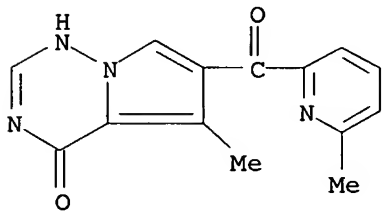
OS MARPAT 139:364963
IT **621685-55-0P 621685-56-1P 621685-58-3P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of aryl ketone pyrrolotriazine compds. as p38 kinase inhibitors)
RN 621685-55-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxaldehyde, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



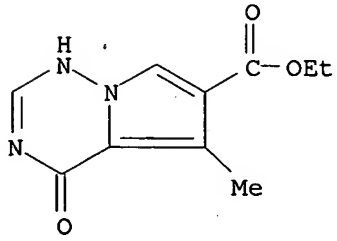
RN 621685-56-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-benzoyl-5-methyl- (9CI) (CA INDEX NAME)



RN 621685-58-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl-6-[(6-methyl-2-pyridinyl)carbonyl]- (9CI) (CA INDEX NAME)



IT **427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(starting material; preparation of aryl ketone pyrrolotriazine compds. as p38 kinase inhibitors)
RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)

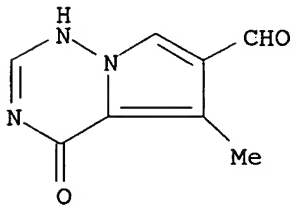


RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

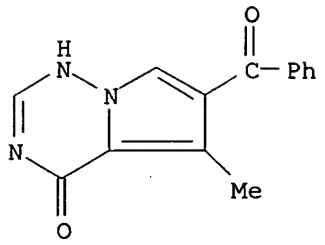
L4 ANSWER 11 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:875173 CAPLUS
DN 139:381511
TI Pyrrolotriazine aniline compounds useful as kinase inhibitors, particularly p38 kinases, and their preparation, pharmaceutical compositions, and use as antiinflammatory agents
IN Dyckman, Alaric; Hynes, John; Leftheris, Katherina; Liu, Chunjian; Wrobleski, Stephen T.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 158 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003090912	A1	20031106	WO 2003-US12426	20030415
	WO 2003090912	C2	20040108		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2483164	AA	20031106	CA 2003-2483164	20030415
	EP 1497019	A1	20050119	EP 2003-724157	20030415
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	BR 2003009669	A	20050301	BR 2003-9669	20030415
PRAI	US 2002-374938P	P	20020423		
	WO 2003-US12426	W	20030415		

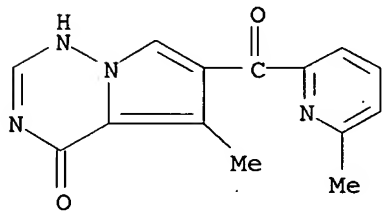
OS MARPAT 139:381511
IT **621685-55-0P 621685-56-1P 621685-58-3P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of pyrrolotriazine aniline compds. as p38 kinase inhibitors)
RN 621685-55-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxaldehyde, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



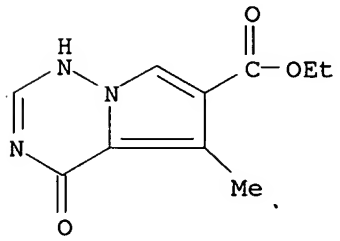
RN 621685-56-1 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 6-benzoyl-5-methyl- (9CI) (CA INDEX NAME)



RN 621685-58-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazin-4(1H)-one, 5-methyl-6-[(6-methyl-2-pyridinyl)carbonyl]- (9CI) (CA INDEX NAME)



IT **427878-70-4**
RL: RCT (Reactant); RACT (Reactant or reagent)
(starting material; preparation of pyrrolotriazine aniline compds. as p38 kinase inhibitors)
RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 12 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:777390 CAPLUS
DN 139:292275
TI Methods for the preparation of pyrrolotriazine compounds useful as kinase inhibitors
IN Godfrey, Jollie Duaine; Hynes, John; Dyckman, Alaric J.; Leftheris, Katerina; Shi, Zhongping; Wrobleski, Stephen T.; Doubleday, Wendel William; Grosso, John A.
PA Bristol-Myers Squibb Company, USA
SO U.S. Pat. Appl. Publ., 36 pp., Cont.-in-part of U.S. Ser. No. 36,293.

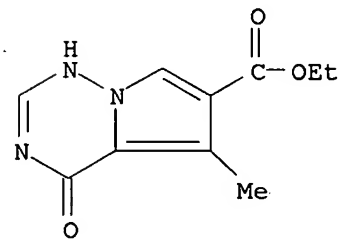
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 2003186982	A1	20031002	US 2002-289010	20021106
US 6867300	B2	20050315		
US 2003069244	A1	20030410	US 2001-36293	20011107
US 6670357	B2	20031230		
ZA 2003003786	A	20040816	ZA 2003-3786	20030515
US 2004229877	A1	20041118	US 2003-696178	20031029
WO 2004043912	A2	20040527	WO 2003-US35220	20031103
WO 2004043912	A3	20040701		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2005107462	A1	20050519	US 2004-19788	20041222
PRAI US 2000-249877P	P	20001117		
US 2001-310561P	P	20010807		
US 2001-36293	A2	20011107		
US 2002-289010	A	20021106		

OS MARPAT 139:292275
IT **427878-70-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolotriazine derivative as kinase inhibitor)
RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 13 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:396849 CAPLUS
DN 138:401758
TI Preparation of 5-substituted N-(1H-indazol-5-yl)pyrrolo[2,1-f][1,2,4]triazin-4-amines as antiproliferative agents
IN Mastalerz, Harold; Zhang, Guifen; Tarrant, James G.; Vite, Gregory D.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 74 pp.
CODEN: PIXXD2
DT Patent

LA English
FAN.CNT 1

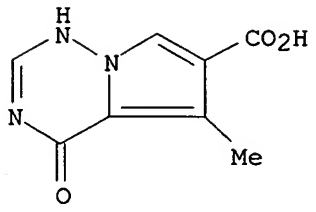
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003042172	A2	20030522	WO 2002-US36528	20021112
	WO 2003042172	A3	20040129		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2467068	AA	20030522	CA 2002-2467068	20021112
	EP 1446401	A2	20040818	EP 2002-793930	20021112
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
	BR 2002014112	A	20040914	BR 2002-14112	20021112
	NZ 533034	A	20041126	NZ 2002-533034	20021112
	JP 2005509030	T2	20050407	JP 2003-544009	20021112
	US 2003186983	A1	20031002	US 2002-294281	20021114
	US 6908916	B2	20050621		
PRAI	US 2001-333014P	P	20011114		
	WO 2002-US36528	W	20021112		

OS MARPAT 138:401758

IT **310435-15-5P**, 5-Methyl-4-oxo-3,4-dihydropyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of N-(indazolyl)pyrrolotriazinamines as tyrosine kinase inhibitors for treatment of proliferative disorders and other diseases associated with signal transduction pathways)

RN 310435-15-5 CAPLUS

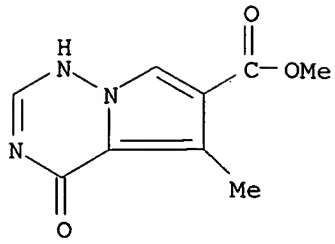
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



IT **310431-29-9**, 5-Methyl-4-oxo-3,4-dihydropyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid methyl ester
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of N-(indazolyl)pyrrolotriazinamines as tyrosine kinase inhibitors for treatment of proliferative disorders and other diseases associated with signal transduction pathways)

RN 310431-29-9 CAPLUS

CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)

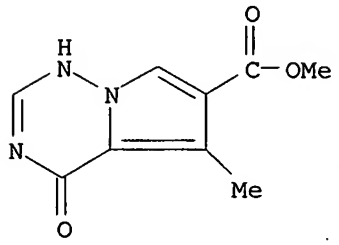


L4 ANSWER 14 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2002:391720 CAPLUS
DN 136:386144
TI Preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivatives for use in treating p38 kinase-associated conditions
IN Leftheris, Katerina; Barrish, Joel; Hynes, John; Wrobleski, Stephen T.
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 108 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2

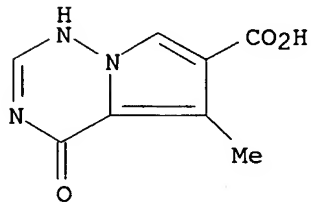
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002040486	A2	20020523	WO 2001-US49982	20011107
	WO 2002040486	A3	20030912		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	CA 2429628	AA	20020523	CA 2001-2429628	20011107
	AU 2002032760	A5	20020527	AU 2002-32760	20011107
	EE 200300227	A	20031015	EE 2003-227	20011107
	EP 1363910	A2	20031126	EP 2001-992298	20011107
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	JP 2004522713	T2	20040729	JP 2002-543494	20011107
	BG 107750	A	20040130	BG 2003-107750	20030421
	ZA 2003003786	A	20040816	ZA 2003-3786	20030515
	NO 2003002229	A	20030716	NO 2003-2229	20030516
PRAI	US 2000-249877P	P	20001117		
	US 2001-310561P	P	20010807		
	WO 2001-US49982	W	20011107		

OS MARPAT 136:386144
IT **310431-29-9P 310435-15-5P 310443-54-0P,** 4-Hydroxy-5-methoxypyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid ethyl ester **427878-70-4P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(intermediate; preparation of pyrrolo[2,1-f][1,2,4]triazine carboxylic acid derivs. for use in treating p38 kinase-associated conditions)
RN 310431-29-9 CAPLUS

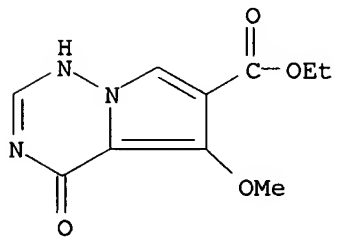
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



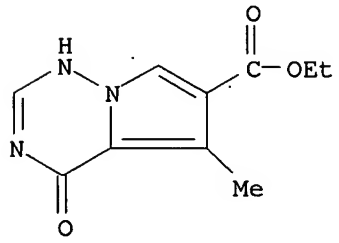
RN 310435-15-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



RN 310443-54-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methoxy-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RN 427878-70-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



L4 ANSWER 15 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2000:841986 CAPLUS
DN 134:17506
TI Preparation of pyrrolotriazines as kinases inhibitors for treating inflammation, cancer, and proliferative diseases
IN Hunt, John T.; Bhide, Rajeev S.; Borzilleri, Robert M.; Qian, Ligang
PA Bristol-Myers Squibb Company, USA
SO PCT Int. Appl., 130 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000071129	A1	20001130	WO 2000-US13420	20000516
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	CA 2373990	AA	20001130	CA 2000-2373990	20000516
	EP 1183033	A1	20020306	EP 2000-930761	20000516
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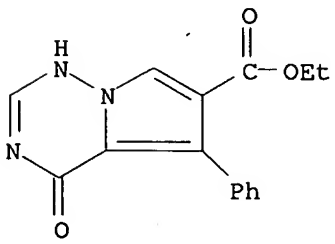
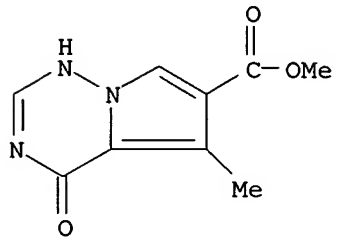
	BR 2000010482	A	20020423	BR 2000-10482	20000516
	JP 2003500359	T2	20030107	JP 2000-619433	20000516
	NZ 516292	A	20040130	NZ 2000-516292	20000516
	AU 770377	B2	20040219	AU 2000-48524	20000516
	TR 200103352	T2	20050321	TR 2001-200103352	20000516
	NO 2001005650	A	20011120	NO 2001-5650	20011120
	ZA 2001009577	A	20030220	ZA 2001-9577	20011120
PRAI	US 1999-135265P	P	19990521		
	US 2000-193727P	P	20000331		
	WO 2000-US13420	W	20000516		

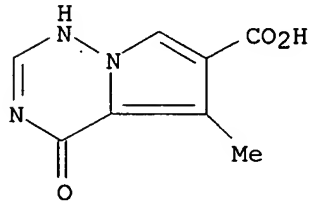
OS MARPAT 134:17506
IT **310431-16-4P 310431-29-9P 310435-15-5P 310436-48-7P 310436-60-3P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)
RN 310431-16-4 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-4-oxo-5-phenyl-, ethyl ester (9CI) (CA INDEX NAME)



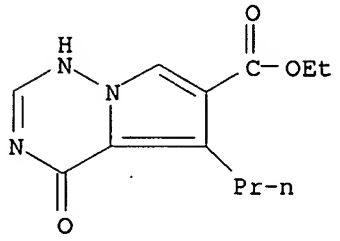
RN 310431-29-9 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo-, methyl ester (9CI) (CA INDEX NAME)



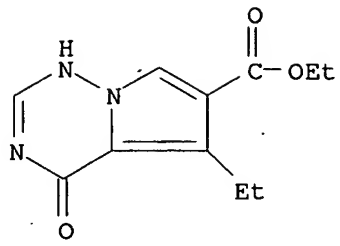
RN 310435-15-5 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methyl-4-oxo- (9CI) (CA INDEX NAME)



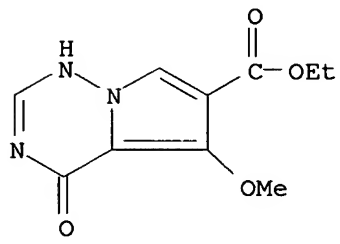
RN 310436-48-7 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-4-oxo-5-propyl-, ethyl ester (9CI) (CA INDEX NAME)



RN 310436-60-3 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 5-ethyl-1,4-dihydro-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



IT **310443-54-0P**
RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of pyrrolotriazines as kinases inhibitors useful in treating inflammation, cancer, and proliferative diseases)
RN 310443-54-0 CAPLUS
CN Pyrrolo[2,1-f][1,2,4]triazine-6-carboxylic acid, 1,4-dihydro-5-methoxy-4-oxo-, ethyl ester (9CI) (CA INDEX NAME)



RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	51.30	212.84

STN INTERNATIONAL LOGOFF AT 18:00:44 ON 23 JUL 2005

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	344	(544/183).CCLS.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:16
L2	40	Bang-Chi.inv. and Chen.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:16
L3	28	Rulin.inv. and Zhao.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:17
L4	79	Joseph.inv. and Sundeen.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:17
L5	57	katerina.inv. and Leftheris.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:18
L6	68	John.inv. and Hynes.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:18
L7	0	Stephen.inv. and Wrobieski.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:19
L8	25	Stephen.inv. and Wrobleski.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:19
L9	7	1 and (2 or 3 or 4 or 5 or 6 or 8)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:20

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	344	(544/183).CCLS.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:16
L2	40	Bang-Chi.inv. and Chen.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:16
L3	28	Rulin.inv. and Zhao.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:17
L4	79	Joseph.inv. and Sundeen.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:17
L5	57	katerina.inv. and Leftheris.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:18
L6	68	John.inv. and Hynes.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:18
L7	0	Stephen.inv. and Wrobieski.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:19
L8	25	Stephen.inv. and Wrobleski.inv.	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:19
L9	7	1 and (2 or 3 or 4 or 5 or 6 or 8)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	OFF	2005/07/23 19:20